

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

I, KAREN BELL, B.A., declare

1. That I am a citizen of the United Kingdom of Great Britain and Northern Ireland, residing at 43 Charingworth Drive, Hatton Park, Warwick, CV35 7SY.
2. That I am well acquainted with the French and English languages.
3. That the attached is a true and correct translation into the English language of the certified copy of French Patent Application No. 0310258 filed on 28th August 2003 and of the official certificate attached thereto.
4. That all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that wilful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such wilful false statements may jeopardise the validity of the patent application in the United States of America or any patent issuing thereon.

Declared this 21st day of May 2010.



KAREN BELL

REPUBLIC OF FRANCE

NIIP

NATIONAL INSTITUTE OF INDUSTRIAL PROPERTY

PATENT OF INVENTION

UTILITY CERTIFICATE

OFFICIAL COPY

The Director-General of the National Institute of Industrial Property certifies that the attached document is the certified true copy of an application for an industrial property title filed at the Institute.

Drawn up in Paris on 25th MAY 2010

On behalf of the Director-General
of the National Institute of
Industrial Property
The Head of the Patent Department

(signature)

Martine PLANCHE

NATIONAL INSTITUTE OF INDUSTRIAL PROPERTY
 26 bis, rue de Saint Pétersbourg 75800 Paris Cedex 08
 For Information: NIIP Direct
 INDIGO No. 0 825 83 85 87
 0.15€ TTC/min
 Fax: 33 1 53 04 52 65

PATENT, UTILITY CERTIFICATE
 Code of intellectual property, Book VI

REQUEST FOR GRANT
 Page 1/2

This printed form is to be legibly completed in black ink

Reserved for the NIIP SUBMISSION OF DOCUMENTS DATE 28 AUGUST 2003 LOCATION 75 NIIP PARIS NATIONAL REGISTRATION NO. ALLOCATED BY THE NIIP 0310258 DATE OF FILING ALLOCATED BY THE NIIP 28 AUGUST 2003	1. NAME AND ADDRESS OF THE APPLICANT OR AGENT TO WHOM CORRESPONDENCE SHOULD BE SENT CABINET LAVOIX 2, Place d'Estienne d'Orves 75441 PARIS CEDEX 09
Your references for this file (optional) BFF 03P0100	
Confirmation of filing by fax	<input type="checkbox"/> Number allocated to the fax copy by the NIIP
2. NATURE OF THE REQUEST Patent application Utility certificate application Divisional application Initial patent application or initial utility certificate application Conversion of a European patent application Initial patent application	Tick one of the following 4 boxes <input checked="" type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> No. Date No. Date <input type="checkbox"/> No. Date
3. TITLE OF INVENTION (max. 200 characters or spaces)	

Membranes for electron conduction and ionic oxygen comprising a layer of mixed vanadium and magnesium oxide

4. DECLARATION OF PRIORITY OR PETITION FOR BENEFIT OF THE FILING DATE OF A PRIOR FRENCH APPLICATION

Country or organisation
Date No.
Country or organisation
Date No.
Country or organisation
Date No.

☐ If there are other priorities, tick the box and use the printed continuation sheet

5. APPLICANT (Tick one of the two boxes)

☒ Artificial person
☐ Physical person

Surname
or name

CENTRE NATIONAL DE LA
RECHERCHE SCIENTIFIQUE
(C.N.R.S)

First names
Legal form

SIREN No
APE-NAF code

Address or headquarters
Road
Postal code and town

3, rue Michel Ange,
75016 PARIS

Country
Nationality
Telephone number (optional)

FRANCE
French

Email address (optional)

Fax number (optional)

☐ If there is more than one applicant, tick the box and use the printed continuation sheet

The second page must be filled in

PATENT, UTILITY CERTIFICATE
REQUEST FOR GRANT
Page 2/2

<p>Reserved for the NIIP</p> <p>SUBMISSION OF DOCUMENTS</p> <p>DATE 28 AUGUST 2003 LOCATION 75 NIIP PARIS</p> <p>NATIONAL REGISTRATION NO. ALLOCATED BY THE NIIP 0310258</p>	
<p>6 AGENT (if applicable)</p> <p>Name First name Office or Company Nationality No. of permanent power of attorney and/or contractual band</p> <p>Address Road Postal code and town Country</p> <p>Telephone number (optional) Fax number (optional) Email address (optional)</p>	<p>CABINET LAVOIX</p> <p>2, Place d'Estienne d'Orves 75441 PARIS CEDEX 09 FRANCE</p> <p>01 53 20 14 20 01 48 74 54 56 brevets@cabinet-lavoix.com</p>
<p>7 INVENTOR(S) The inventors are the applicants</p>	<p>The inventors are physical persons <input type="checkbox"/> yes <input checked="" type="checkbox"/> no If answer is no, provide separate designation of inventor(s)</p>

<p>8 SEARCH REPORT</p> <p>Immediate drawing-up or deferred drawing-up</p> <p>Payment of fee in instalments (in two transactions)</p>	<p>Only for a patent application (including Divisional and Conversion)</p> <p><input checked="" type="checkbox"/> <input type="checkbox"/></p> <p>Only for physical persons carrying out their own filing <input type="checkbox"/> yes <input type="checkbox"/> no</p>
<p>9 REDUCTION OF FEE RATE</p>	<p>Only for physical persons <input type="checkbox"/> requested for the first time for this invention (attach notice of non-imposition)</p> <p><input type="checkbox"/> obtained prior to this filing for this invention (attach a copy of the decision to allow free aid or indicate its reference) AG</p>
<p>10 SEQUENCES OF NUCLEOTIDES AND/OR AMINO ACIDS</p> <p>The electronic data medium is attached</p> <p>The declaration of conformity of the sequence listing in paper form with the electronic data medium is attached</p> <p>If you have used the printed continuation sheet, indicate the number of pages attached</p>	<p><input type="checkbox"/> Tick the box if the description contains a sequence listing</p> <p><input type="checkbox"/></p> <p><input type="checkbox"/></p>
<p>11 SIGNATURE OF THE APPLICANT OR OF THE AGENT (name and capacity of signatory)</p> <p>B. DOMENEGO No. 00-0500 (Signature)</p>	<p>VISA OF PREFECTURE OR NIIP (Signature)</p>

Law No. 78-17 of 6 January 1978 relating to data processing, data files and freedoms applies to the replies made on this form. It guarantees a right of access and correction in respect of data concerning you at the NIIP.

NATIONAL INSTITUTE OF INDUSTRIAL PROPERTY
26 bis, rue de Saint Pétersbourg 75800 Paris Cedex 08
For information: NIIP DIRECT
Indigo No. 0 825 83 85 87
0.15€ TTC/min.
Fax: 33 1 53 04 52 65

PATENT, UTILITY CERTIFICATE
Code of intellectual property, Book VI

DESIGNATION OF INVENTOR(S)

Page No. 1/2

(to be provided if the applicants are not the inventors)

This printed form is to be legibly completed in black ink

Your references for this file (optional)	BFF 03P0100
NATIONAL REGISTRATION NO.	03 10258
TITLE OF INVENTION (max. 200 characters or spaces) Membranes for electron conduction and ionic oxygen comprising a layer of mixed vanadium and magnesium oxide	
THE APPLICANT(S): CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE (C.N.R.S)	
DESIGNATE(S) AS INVENTORS:	
1. Surname First names Road Address Town and post code Employing company (optional)	FARRUSSENG David 22, rue Pierre Brunier 69300 CALUIRE FRANCE
2. Surname First names Road Address Town and post code Employing company (optional)	MIRODATOS Claude 17, rue Montaigne 69003 LYON FRANCE

3. Surname First names Road Address Town and post code Employing company (optional)	REBEILLEAU Michael 96, rue Montesquieu 69007 LYON FRANCE
If there are more than three inventors, use more than one form. Indicate at the top-right hand side the page number followed by the number of pages.	
DATE AND SIGNATURE(S) OF APPLICANT(S) OR (Signature) REPRESENTATIVE (name and capacity of signatory) Paris, 4 th March 2004 L. BOUGET No. 92-1033	

Law No. 78-17 of 6 January 1978 relating to data processing, data files and freedoms applies to the replies made on this form. It guarantees a right of access and correction in respect of data concerning you at the NIIP.

NATIONAL INSTITUTE OF INDUSTRIAL PROPERTY
 26 bis, rue de Saint Pétersbourg 75800 Paris Cedex 08
 For information: NIIP DIRECT
 Indigo No. 0 825 83 85 87
 0.15€ TTC/min.
 Fax: 33 1 53 04 52 65

PATENT, UTILITY CERTIFICATE
 Code of intellectual property, Book VI

DESIGNATION OF INVENTOR(S)

Page No. 2/2

(to be provided if the applicants are not the inventors)

This printed form is to be legibly completed in black ink

Your references for this file (optional)	BFF 03P0100
NATIONAL REGISTRATION NO.	03 10258
TITLE OF INVENTION (max. 200 characters or spaces) Membranes for electron conduction and ionic oxygen comprising a layer of mixed vanadium and magnesium oxide	
THE APPLICANT(S): CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE (C.N.R.S)	
DESIGNATE(S) AS INVENTORS:	
1. Surname First names Road Address Town and post code Employing company (optional)	VAN VEEN André 36, rue des Antonins 69100 VILLEURBANNE FRANCE
2. Surname First names Road Address Town and post code Employing company (optional)	RUSHWORTH Simon 12, Mayew Road Irby Wirral CH61 3XR GREAT BRITAIN

3. Surname First names Road Address Town and post code Employing company (optional)	ROUSSET Jean Luc 16, rue de Créqui 69006 LYON CEDX 06 FRANCE
If there are more than three inventors, use more than one form. Indicate at the top-right hand side the page number followed by the number of pages.	
DATE AND SIGNATURE(S) OF APPLICANT(S) OR (Signature) REPRESENTATIVE (name and capacity of signatory) Paris, 4 th March 2004 L. BOUGET No. 92-1033	

Law No. 78-17 of 6 January 1978 relating to data processing, data files and freedoms applies to the replies made on this form. It guarantees a right of access and correction in respect of data concerning you at the NIIP.

The invention relates to membranes which have electron and oxygen ion conducting properties, the method of preparation thereof and the applications thereof, in particular for the oxidative dehydrogenation of alkanes into alkenes.

Oxidative dehydrogenation methods which allow saturated organic compounds to be converted into unsaturated compounds are well known. A large number of attempts to improve methods of this type have been made over recent years in order to increase in particular the conversion rates and the selectivity with regard to the desired products.

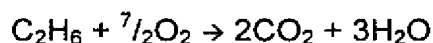
Ethylene is produced industrially by means of thermal cracking of ethane, generally in the presence of water vapour, which consists in a pyrogenic reaction of ethane at high temperatures in the order of 850°C. At the present time, the selectivity rate in terms of ethylene obtained according to this method is not higher than 80% and the yield in terms of ethylene is in the order of 60%. Although an increase in the temperature allows higher conversion rates to be achieved, this increase, however, brings about a reduction of the selectivity by promoting the formation of secondary products and their decomposition in the form of coke. The depositing of coke on the walls of the pipes of the installation principally brings about two harmful effects; on the one hand, it results in pressure drops through the pipes of the installation and, on the other hand, it offers resistance to the transfer of heat to the hydrocarbon fluid.

The formation of coke constitutes one of the main factors limiting cracking operations.

In order to attempt to overcome this limitation, methods for oxidative dehydrogenation of ethane have been developed.

There are substantially two different types of method for oxidative dehydrogenation: catalytic methods which use catalyst beds and those which use mixed conducting membranes.

The main reactions which are involved in the catalytic method are as follows:



Catalysts which allow yields for converting ethane into ethylene in the order of 45-50% to be achieved at temperatures in particular less than 700°C have recently been described.

In this manner, Thorsteinson et al. (*J. Catal.*, 52, 1978, 116) obtained an ethylene yield of 25% with a selectivity of 90% using the catalyst MoVNbTeOx at 350°C.

Higher yields of between 44 and 50% have also been reported by Ji and Liu et al. (L. Ji, J. Liu, *Chem. Commun.*, 1996, 1203) and Velle et al. (O.J. Velle, A. Andersen, K.-J. Jens, *Catal. Today*, 6 (1990) 567) with the catalysts La/CaO, Li/La/CaO and $\text{SrCe}_{0.5}\text{Yb}_{0.5}\text{O}_{2.75}$ in a fixed bed reactor, but at temperatures in the order of from 600 to 700°C.

However, in order to achieve a yield of this type, it is necessary to operate in an oxygen-enriched atmosphere, which brings about an increase in operating costs.

Generally, although they allow the problem of coke formation to be overcome, catalytic methods for oxidative dehydrogenation are nonetheless difficult to use on an industrial scale at the present time. They have a number of disadvantages, in particular yields which are lower than those obtained using commercial methods, the consumption of the hydrogen produced, the formation of oxygenated secondary products, the inflammable nature of the reaction mixture, the requirement for oxygen enrichment and the costs which result.

The methods for selective oxidative dehydrogenation of ethane into ethylene which use dense oxygen ion conducting membranes have been found to be much more promising.

Dense conducting membranes are membranes which are capable of selectively transporting oxygen ions at temperatures which are generally higher than 600°C, most often between 700°C and 1200°C.

The different types of dense membrane for conducting oxygen include in particular membranes composed of solid electrolyte, the ion type, and mixed ion-electron conducting membranes based on multimetal oxides.

Membranes which are formed from solid electrolyte comprise inorganic oxides, typically oxides of calcium or zirconium, which are stabilised by yttrium (YSZ) and generally have a perovskite or fluorite structure.

These membranes of the ion type transport only oxygen ions and require the application of an external electric field in order to maintain an electron flow through the membrane and thus the process of ionisation-deionisation.

Mixed ion-electron conducting membranes, in particular monophase membranes which are constituted by multimetal oxides, are, however, capable of transporting both oxygen ions and electrons without it being necessary to apply an external electric field.

The driving force which allows oxygen to be transported in mixed conducting membranes is based on a partial pressure difference of O_2 applied at one side and the other of the membrane. Owing to its non-porous structure, the membrane prevents any gas molecule from passing through directly. Only the oxygen ions are able to selectively migrate. The dissociation and ionisation of the oxygen is brought about in contact with the surface where the partial pressure is highest (cathode), by means of electron capture. The charge flow of the oxygen ions O^{2-} is compensated for by a simultaneous flow of electrons in the opposite direction. When the oxygen ions reach the side of low partial oxygen pressure, that is to say, the side of the permeate (anode), the oxygen ions donate their electrons and recombine in order to regenerate molecular oxygen which is released into the permeation current.

These membranes, which are used as membrane reactors for the oxidative dehydrogenation of ethane, selectively transport oxygen as far as the anode where they react rapidly with ethane in order to form ethylene.

The transport of oxygen through these conduction membranes is a controlled process which is substantially dependent on two factors: the kinetic surface limitations and the limitations of volume diffusion.

The kinetic surface limitations are linked to the plurality of steps which are involved during the conversion of the oxygen molecules in a gaseous phase in the charge current into mobile oxygen ions and, conversely, of the mobile oxygen ions into oxygen molecules at the side of the permeate of the conducting membrane or during the reaction of the oxygen ions with the reactant gas.

The limitations of volume diffusion are linked to the diffusion of the oxygen ions and electrons at the inner side of the membrane.

These two limiting factors are dependent in particular on the partial pressure gradient in terms of O_2 at one side and the other of the membrane, the operating temperature and the thickness of the membrane.

Work has been carried out which is intended to optimise the efficiency levels of these membranes, which are advantageous in particular when used as membrane reactors for the oxidative dehydrogenation of ethane into ethylene.

A first approach involves seeking multimetal mixed conducting oxide compositions which have good intrinsic properties for transporting oxygen.

Document Haihui Wang et al., Chem. Comm., 2002, 1468-1469, describes a reaction for oxidative dehydrogenation of ethane into ethylene in a membrane reactor which is constituted by $Ba_{0.5} Sr_{0.5} Co_{0.8} Fe_{0.2} O_{3-\delta}$.

The ethylene is obtained at a yield of 15%, a conversion rate of 18%, a selectivity of 90% at 650°C and an oxygen permeation flow of 0.36 mL.min⁻¹ cm⁻².

In the document H. Wang et al. (*Catalysis Letters*, 2002, vol. 84, Nos 1-2 pp. 101-106), the oxidative dehydrogenation reaction of ethane in the Ba_{0.5}Sr_{0.5}Co_{0.8}Fe_{0.2}O_{3-δ} membrane reactor is examined at 800°C. A maximum yield of 67% is obtained with a selectivity with regard to ethylene, a conversion rate and a permeation flow of oxygen of 80%, 82% and 1.6 mL cm⁻² min⁻¹, respectively.

Another approach involves reducing the kinetic surface limitations by associating a catalyst which promotes, for example, the dissociation of gaseous oxygen into mobile O²⁻ ions.

Document EP 0 663 231 (Air Products and Chemicals, Inc.) thus describes a membrane which comprises a porous mixed conducting layer of multimetal oxide, one surface of which layer is covered with a catalyst and the other is in contact with a dense mixed conducting multimetal layer. The catalyst comprises a metal or a metal oxide which catalyses the dissociation of oxygen molecules into ions and/or the association of oxygen ions into molecular oxygen.

The two membranes which are set out by way of example in this patent application use a platinum-based catalyst at a ratio of 10 mg/cm² of surface-area. The use of an expensive precious metal in a substantial quantity as a catalyst renders the use of these membranes on an industrial scale improbable.

Document WO 99/21649 describes a catalytic membrane reactor which comprises an oxidation zone and a reduction zone which are separated by a membrane which is impermeable to gases, and which has an oxidation surface in contact with the oxidation zone and a reduction surface which is in contact with the reduction zone, a layer of adhesive catalyst on the oxidation surface of the membrane and, optionally, a three-dimensional catalyst in the oxidation zone. However, the catalysts are in the form of a continuous coating or layer.

Furthermore, only the catalysts $\text{La}_{0.8} \text{Sr}_{0.2} \text{MnO}_3$ and cermet of Ni on $\text{La}_{0.8} \text{Sr}_{0.2} \text{MnO}_3$ are actually envisaged. There is no mention anywhere of dispersed particles based on magnesium or noble metals.

These membranes are difficult to use and attempts have been made to produce membranes which are effective, less costly and in particular, much easier to use.

Surprisingly, it has now been discovered that, by dispersing a small quantity of catalyst based on magnesium oxide or noble metals in the form of particles at the surface of a mixed conducting membrane of multimetal oxide, it was possible to obtain less costly membranes which have high rates of conversion and selectivity.

More particularly, it has been shown that this modification of the surface of the membrane allows the flow of oxygen and/or the catalytic yield to be significantly increased, and/or the operating temperature conditions to be reduced. By way of example, the operating temperature conditions can be reduced by from 10 to 20%, which may represent approximately a hundred degrees.

This temperature reduction has a number of advantages. In addition to the reduction in the operating cost which a temperature reduction of approximately a hundred degrees represents for industrial production, this allows the problems of corrosion in the region of the reactor to be substantially reduced, and also offers a wider selection of materials for the construction of the reactor itself.

It also allows the use of a larger number of reactions to be envisaged, in particular those which require high temperatures, such as the partial oxidation of light alkanes.

In a particularly advantageous manner, it was found that placing a dense mixed conducting membrane of multimetal oxide in contact with dispersed particles based on magnesium oxide or noble metal modifies the redox properties of the membrane.

Without wishing to be limited to any one theory, it is considered that particles based on magnesium oxide or noble metals dispersed on the oxidation surface (or anode) of the membrane may promote the recombination of oxygen ions into O_2 , then the desorption of the molecules of O_2 .

Furthermore, the presence of the dispersed particles based on magnesium oxide or noble metals on one of the surfaces of the membrane allows the thermal stability of the resulting membrane to be increased. The membranes are thus found to be much more stable, in particular in terms of the mechanical and thermal stresses to which they are subjected under the operating conditions, retaining their structural integrity for a longer period of time.

One of the objects of the present invention is to provide mixed conducting membranes of multimetal oxides which are modified at the surface and which have an improved flow in terms of oxygen in comparison with bare membranes.

According to one advantageous feature of the invention, the composite membranes according to the invention allow an increase of the permeation flow of the oxygen, generally of between 5 and 20%, and advantageously in the order of 30%.

Another object of the present invention is to provide membranes which have high catalytic activity at lower operating temperatures.

In the context of the present description, "high catalytic activity" is intended to refer to a level of activity which allows yields to be reached which are greater than 60%, preferably greater than 70%, and more preferably greater than 80%. By way of illustration, the membranes according to the invention allow ethane to be converted into ethylene with a yield greater than 65%.

Another object of the present invention is to provide a method for preparing these membranes. Advantageously, this is easy to implement, inexpensive and can be readily used on an industrial scale, which should promote the use and the development of the membrane reactors of the invention on an industrial scale.

Another object of the present invention is to provide membrane catalytic reactors which comprise the membranes according to the invention. Advantageously, these membrane

reactors allow high levels of conversion and selectivity to be reached.

The membranes according to the invention are advantageous in a large range of methods, including in particular the separation of oxygen from a gaseous mixture or the oxidation of light hydrocarbons.

Advantageously, the membranes according to the invention have an increased permeation flow of oxygen and therefore allow the oxygen to be extracted from a gaseous mixture with a high yield at a lower temperature in comparison with oxygen conducting membranes which are bare, that is to say, which have no deposit of dispersed particles based on magnesium oxide or noble metals.

In particular, the membranes according to the invention are particularly advantageous when used as a membrane reactor for the oxidative dehydrogenation of ethane, in that they allow levels of conversion, selectivities and yields to be obtained which are higher than those described in the prior art.

Another object of the present invention is therefore to provide a method for oxidative dehydrogenation of ethane into ethylene which constitutes an advantageous alternative to the method for thermal dehydrogenation of ethane.

In addition to the high levels of conversion and selectivities, this method advantageously allows the coking problems of the method for thermal dehydrogenation to be overcome. Furthermore, the separation of the oxygen coupled with the consumption thereof in the oxidative dehydrogenation reaction brings about a number of advantages: the formation

of secondary oxygenated products is limited, air, in particular at atmospheric pressure, can be used directly as a source of oxygen without it being necessary to carry out a cryogenic or other distillation operation, and the operating conditions can be controlled with no danger of explosion for the installation.

These objects and others are achieved by the invention which relates, according to a first aspect, to an oxygen conducting membrane, comprising a dense mixed conducting membrane of multimetal oxide, one of the surfaces of which membrane is coated with dispersed particles based on magnesium oxide or noble metals.

MEMBRANES

Dense mixed conducting membrane of multimetal oxides

In the context of the present description, the term "dense" is intended to refer to a layer which is "impermeable to gaseous fluids", i.e. which does not allow the passage of a significant quantity of a gaseous fluid through the membrane. In specific cases, a low level of permeability in terms of gases other than O_2 may be acceptable or inevitable. This is the case, for example, when the membrane comprises structural defects or when the sealing of the reactor is defective.

Generally, the expression "dense membrane" in the context of the present invention includes membranes which have a gas permselectivity greater than 30, and preferably greater than 100.

Dense mixed conducting membranes of multimetal oxide transport the oxygen anions and are in this respect permeable

to oxygen. On the other hand, they are impermeable to the gaseous oxygen itself.

In the context of the present description, "mixed conducting" is intended to refer to a dense layer which is capable of transporting both electrons and oxygen ions in a selective manner.

The dense layer of the invention is formed from a single multimetall oxide or a mixture of two or more multimetall oxides, each multimetall oxide comprising an oxide of at least two different metals or a mixture of at least two different metal oxides, in which the multimetall oxide has electron and oxygen-ion conductivity at temperatures higher than approximately 500°C. It is also possible to refer to a solid solution of multimetall oxide.

In the context of the present description, "operating temperature" is intended to refer to a temperature at which the membrane according to the invention has a significant level of electron and oxygen-ion conductivity.

Generally, the operating temperatures are between approximately 500°C and 1100°C. The specific temperature, and the flows of reactant gas and gas containing oxygen, are optimised in accordance with the membrane, the catalyst particles and the envisaged application. The temperature must be sufficiently high to promote the oxygen anion flow through the membrane without impairing the membrane and the catalyst particles.

The dense multimetall oxide mixed conducting layer of the invention therefore designates a membrane which has electron

and oxygen-ion conductivity and the capacity to separate the oxygen from a gaseous mixture containing oxygen and one or more other volatile components using the conductivities.

According to a preferred method, the dense layer is formed from mixed oxides which have a perovskite structure.

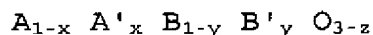
The compounds of the perovskite type comply with the formula ABO_3 , in which A represents an ion of an alkali metal, alkaline-earth metal or rare earth metal and B an ion of a transition metal. Ideally, they have a cubic crystalline structure which is very stable and in which the ions A are positioned at the peaks of the cubic crystalline mesh, the oxygen ions are located at the centre of each face of the cube and form an octahedron BO_6 with B placed at the centre of the cube. Ideal perovskite structures generally have a valency total for the ions A and B equal to 6, as in the mineral perovskite model $CaTiO_3$.

The substitution of a site A or B by another alkaline-earth metal or metal cation can produce sites which are vacant in terms of oxygen. These vacant sites allow the transport of oxygen through the crystalline network.

Examples of perovskite structure are in particular cubic perovskites, Brownmillerites, Aurivillius phases or lamellar perovskites and the like.

Preferably, the perovskite structures in the context of the present description comprise structures which have a perovskite phase greater than 90%, more preferably greater than 95% and even more preferably greater than 98%.

Preferably, the dense mixed conducting membrane refers to a compound having the general formula:



in which

A and A', which may be the same or different, each represent a metal ion or an alkaline-earth metal or a metal which is selected from the lanthanide series;

B and B', which may be the same or different, each represent a metal ion and/or a mixture of metal ions in which the metal is selected from the transition metals;

$$0 \leq x \leq 1;$$

$$0 \leq y \leq 1;$$

z is a number which renders the charge of the compound neutral and which defines the oxygen deficiency; z is thus dependent on the nature of A, A', B, B', x, y and the temperature.

A and A', which may be the same or different, preferably represent La, Ca, Sr, and/or Ba, more preferably Ba and/or Sr.

According to a specific method, it is preferable for A and A' to be different.

It is in particular preferable for A to represent Ba. A' preferably represents Sr.

According to a preferred method, B and B', which may be the same or different, represent Cr, Mn, Fe, Co, Ni and/or Cu, more preferably Co and/or Fe.

B and B' are preferably different.

anisotropic morphology and which may or may not have a crystalline structure, and which have a mean diameter which is less than or equal to $3\mu\text{m}$, preferably with a narrow granulometric distribution.

In the context of the present description, "dispersed" is intended to refer to a distribution of separate particles which are distributed in a random or organised manner and which form a discontinuous deposit at the surface of the membrane, given that the zones covered by the particles represent between 10% and 80% of the surface-area of the membrane.

It should be noted that the zones of the surface of the membrane which is not covered by particles based on magnesium oxide or noble metals are bare, that is to say, they are directly exposed to the reactant gases or the gas which contains oxygen. That is to say, this modified surface comprises no other coating.

The dispersion of particles based on magnesium oxide or noble metals which are directly in contact with a dense membrane allows the formation of "triple points", that is to say, an increase in the contact surface-area between the catalyst, that is to say, the particles based on magnesium oxide or noble metals, the reactant gas and the "active" oxygen (O_2) at the surface of the membrane.

Advantageously, this type of deposit further allows the limitation of reabsorption phenomena of the product formed on the oxidation surface by the catalyst which may lead to the degradation thereof. This therefore advantageously becomes evident in an improvement of the selectivity, in particular

in the case in which the products produced are sensitive to the action of the catalyst, such as, for example, in the case of the oxidative dehydrogenation of ethane into ethylene.

In the context of the present description, the term "particles based on magnesium oxide" generally refers to any particle which is constituted completely or partially by a magnesium oxide, the magnesium oxide at the centre of the material being present in at least one of the following forms:

- in the form of magnesium oxide having the formula MgO ;
and/or
- in the form of magnesium oxide referred to as "doped", that is to say, a magnesium oxide which includes metal cations for insertion or substitution.

In a doped magnesium oxide, the magnesium cations generally remain in the majority, that is to say, the molar ratio of the quantity of magnesium relative to the total quantity of cations in the doped magnesium oxide generally remains greater than 50% by mole, most often greater than 70% and, in most cases, greater than 90% by mole.

The magnesium oxide is preferably doped using a transition metal, preferably selected from vanadium.

According to a particularly preferred aspect, the magnesium oxide is doped using vanadium, preferably using from 0.5 to 10% by weight of vanadium, more preferably using 5% by weight of vanadium.

"Particles of noble metals" are understood to be particles which are constituted substantially by a noble metal selected from Pd, Pt, Rh, Au, Ru, Ag, and Ir or an alloy thereof, the

noble metal being completely or partially in reduced or oxidised form. Amongst the noble metals, Pd is particularly preferred.

The particles based on magnesium oxide or noble metals are preferably dispersed in accordance with a density of between $1 \cdot 10^5$ and $1 \cdot 10^7$ particles per cm^2 and $1 \cdot 10^{12}$ and $5 \cdot 10^{12}$ particles per cm^2 , respectively.

According to a preferred method, the discontinuous deposit formed by the particles based on magnesium oxide or noble metals has a specific surface-area which varies from 30 to 50 m^2/g , preferably between 40 and 50 m^2/g at an operating temperature of 800°C .

Preferably, the particles based on magnesium oxide or noble metals have a mean diameter of between 5 nm and 3000 nm, more preferably between 5 and 50 nm, and even more preferably between 5 and 10 nm.

This mean particle size advantageously allows a large specific surface-area to be made available, that is to say, allows the contact surface between the gas and the modified membrane to be increased per unit of volume.

The percentage by weight of the deposit formed by the particles based on magnesium oxide or noble metals relative to the weight of the dense mixed conducting layer can vary between approximately 0.01 and 1%, preferably between 0.01 and 0.5%, more preferably between 0.01 and 0.1%.

METHOD FOR PREPARING MEMBRANES ACCORDING TO THE INVENTION

Catalyst based on magnesium oxide

According to a second aspect, the invention relates to the methods for preparing the oxygen conducting membranes defined above.

The particles based on magnesium oxide can be dispersed and deposited by chemical means at the surface of the dense layer of multimetal oxide using conventional techniques, such as the depositing of colloidal suspensions, still referred to as the sol-gel method, or depositing in a vapour phase.

According to a specific embodiment, the membranes which are coated with dispersed particles based on magnesium oxide according to the invention are prepared in accordance with the methods referred to as "sol-gel", which are known to the person skilled in the art.

For further details relating to this matter, reference can be made in particular to the thesis of Laure Albaric, University of Montpellier II, 1999.

This method comprises steps which consist in:

- a. providing a mixed conducting membrane of multimetal oxide as defined above;
- b. preparing a colloidal suspension based on magnesium oxide in an organic solvent;
- c. placing the suspension obtained in contact with the dense mixed conducting membrane; and
- d. calcining the membrane obtained.

The colloidal suspension based on magnesium oxide can be prepared according to the methods referred to as "sol-gel" methods, which are known to the person skilled in the art.

The preparation of colloidal suspension thus comprises the dissolution of the precursor of the particles based on magnesium oxide in the organic solvent, in the desired proportions. The concentration of the precursor solution is fixed in accordance with the coating which it is desirable to obtain and the agitation is generally carried out at ambient temperature until a colloidal suspension is obtained.

The precursors are preferably distilled before they are used so as to eliminate possible traces of water and condensed matter.

By way of example of precursors of particles based on magnesium oxide, it is possible to mention in particular alkoxides, hydroxides, nitrates, magnesium alkoxides, such as magnesium dimethoxide.

According to a preferred embodiment, the magnesium oxide is doped using vanadium. In this case, the colloidal suspension is prepared from an admixture of precursors based on magnesium and vanadium in the desired proportions. Preferably, a solution of magnesium alkoxide and vanadium alkoxide is prepared, preferably of magnesium dimethoxide and vanadium isopropoxide.

The organic solvents used during step b) are those which are commonly used for the preparation of colloidal suspensions and include in particular methanol, ethanol, 2-propanol.

The colloidal suspension produced in step b) is then deposited at the surface of the dense mixed conducting membrane according to the known technique of "spin-coating". In this regard, reference can be made in particular to Hiromichi Sakamoto, Jianbei Qiu and Akio Makishima, *Science and Technology of Advanced Materials*, Volume 4, Issue 1, 2003, Pages 69-76. This technique generally consists in depositing a few millimetres of colloidal suspension, for example, using a syringe, at the surface of the dense membrane, and distributing the colloidal suspension in a homogeneous manner on the surface of the membrane, placing the membrane on a support which is rotated at high speed.

The deposit is then dried in air, then in an oven at 100°C for a length of time sufficient to eliminate the solvent.

The resulting membrane is then calcined at a temperature which is sufficient to eliminate the organic residues, generally at a temperature higher than 400°C, preferably between 800°C and 900°C, and more preferably between 800°C and 850°C, by means of which a membrane is obtained which is coated with dispersed particles based on magnesium oxide.

The dense mixed conducting membrane used according to this method can be prepared according to conventional methods which generally comprise:

- dissolving in water metal salts which constitute the mixed conducting oxide, including in particular nitrates and chlorides, in the desired stoichiometric proportions;
- co-precipitating the salts from the aqueous solution by means of the addition of an organic acid, for example, citric acid;

- isolating the co-precipitate by means of filtration or centrifuging;
- calcining the co-precipitate in order to obtain a multimetal mixed conducting oxide;
- grinding the mixed conducting oxide into the form of powder;
- milling the mixed conducting oxide powder, optionally in association with additives, into the desired form; and
- sintering the milled powder in order to obtain a dense ceramic mixed conducting material.

These methods, which are well known to a person skilled in the art, are suitable for the preparation of the dense mixed conducting membrane of the membranes according to the invention.

The "sol-gel" method is particularly suitable for obtaining particles of micrometric or submicrometric size and dispersing them at the surface of the membrane. The size of the particles formed can be evaluated after the particles have been deposited at the surface of the membrane by means of scanning electron microscopy and/or physisorption.

Catalyst based on noble metals

Membranes which are coated with dispersed particles of noble metals according to the invention can be prepared according to the method which consists in:

- a) providing a mixed conducting membrane of multimetal oxide as defined above;
- b) depositing the noble metal particles by means of laser vaporisation.

The principle of the technique of laser vaporisation is set out in particular in the bibliographic reference "Ion-scattering study and Monte Carlo simulations of surface segregation in Pd-Pt nanoclusters obtained by laser vaporization of the bulk alloys", J.-L. Rousset et al., *Physical Review B*, 15 July 1998, II, vol. 58, N° 4.

Advantageously, this technique allows small metal particles to be deposited with a narrow and homogeneous distribution, a controlled composition and quantity.

The size and the dispersion or density of the particles on the surface of the dense membrane can be determined by means of transmission electron microscopy and/or physisorption.

According to a third aspect, the invention relates to membranes which can be produced according to one of the methods defined above.

The membranes according to the invention can be prepared in different forms, suitable for a specific model of reactor, including in particular in the form of discs or tubes.

The mixed conducting membranes according to the invention are advantageously stable at temperatures of between 25°C and 1100°C.

MEMBRANE REACTOR

According to a fourth aspect, the invention also relates to a catalytic membrane reactor which comprises an oxidation zone and a reduction zone which are separated by a membrane according to the invention.

In the following, the oxidation surface (anode) designates the surface which is in contact with the oxidation zone and the reduction surface (cathode), the one in contact with the reduction zone.

The oxidation surface preferably corresponds to that which is in contact with the dispersed particles based on magnesium oxide or noble metals.

The reduction surface is in contact with the gaseous mixture which contains oxygen. It preferably corresponds to the dense layer, which is optionally covered with a catalytic layer, and is the seat for the dissociation reactions of the molecular oxygen into oxygen ions O^{2-} .

This reduction surface can be optionally coated with a catalyst which promotes the reduction of matter containing oxygen, such as O_2 , NO_2 , SO_2 , in order to produce oxygen anions in the region of the membrane. By way of example of catalysts of this type, it is possible to mention in particular noble metals.

APPLICATION OF THE MEMBRANE REACTOR

The membranes according to the invention are advantageous when used as membrane reactors for the partial or total oxidation of reduced matter, in particular hydrocarbons.

According to a fifth aspect, the invention relates to a method for oxidising a reactant gas comprising:

- i) using a membrane reactor according to the invention;
- ii) introducing reactant gas into the oxidation zone;

- iii) introducing gas containing oxygen into the reduction zone;
- iv) heating the membrane which separates the oxidation and reduction zones at an operating temperature in order to reduce the gas containing oxygen, transport the oxygen ions to the oxidation zone and oxidise the reactant gas.

The operating temperature is preferably between 500 and 800°C.

This method is preferably used for a partial oxidation of the reactant gas, such as a conversion of methane into carbon monoxide and hydrogen, or an oxidative dehydrogenation of hydrocarbons, or a controlled oxidation of hydrocarbons into oxygenated molecules.

According to a preferred embodiment, the reactant gas is a light hydrocarbon which is oxidised into alkene.

In the context of the present description, "light hydrocarbon" is intended to refer to volatile alkanes which contain from 1 to 6 atoms of carbon, including in particular methane, ethane, propane and the butanes.

Light hydrocarbon is preferably ethane or propane, preferably ethane which is oxidised into ethylene.

According to another advantageous feature, the membranes according to the invention have a level of activity for the dehydrogenation of ethane into ethylene greater than that obtained with the dense membrane which has an oxidation surface which is not modified by dispersed particles based on magnesium oxide or noble metals.

According to another aspect, the invention relates to the use of membrane reactors in order to recover oxygen from a gaseous mixture which contains oxygen.

This use comprises in particular the steps which consist in:

- introducing a gaseous mixture containing oxygen into the reduction zone which is separated from the oxidation zone by the membrane according to the invention, by establishing a positive partial pressure difference between the two zones by producing an excess partial pressure of oxygen in the first compartment and/or by producing a reduced partial pressure in the second compartment;
- placing the mixture containing the oxygen in contact with the reduction surface, at a temperature greater than approximately 500°C in order to convert the gaseous mixture containing oxygen into an oxygen enriched current (permeate) on the one hand, and, on the other hand, into an oxygen-depleted current;
- recovering the oxygen enriched current (permeate).

The expression "gaseous mixture containing oxygen" generally refers to gases and gaseous mixtures, one of the components of which is oxygen or an oxide. These can be reduced on the reduction surface of the membrane according to the invention. This expression includes in particular oxides of carbon, nitrogen and sulphur (CO_x , NO_x and SO_x) and gaseous mixtures which comprise an oxide in the presence of an inert gas or any other gas which is non-reactive relative to the membrane.

The expression also includes mixtures of molecular oxygen (O_2) in other gases, for example, oxygen of atmospheric air, oxygen in the presence of an inert gas, such as He, Ar, etcetera.

Legend: Y ethy, Y propy represent the yields in terms of ethylene and propylene, respectively.

Figure 11: Yield from the oxidative dehydrogenation reaction in accordance with temperature using a membrane according to example 3. Conditions: Reduction zone, $F_{AIR}=50\text{mL/min}$, $P_{AIR}=1.2\text{bar}$; oxidation zone, $F_{C_3H_8/He}=37\text{mL/min}$, $P_{C_3H_8}=0.20\text{ atm}$.

Figure 12: Illustration of the surface of the membrane coated with dispersed particles of VMgO according to example 1 using electron scanning microscopy (Scale: $10\mu\text{m/cm}$).

Figure 13: Illustration of the surface of the membrane coated with dispersed particles of Pd according to example 3 using electron transmission microscopy (Scale: $25\mu\text{m/cm}$).

EXAMPLES

Abbreviations

HC: hydrocarbon

He: helium

C_2H_6 : ethane

C_2H_4 : ethylene

C_3H_8 propane

F_{AIR} , $F_{C_2H_6}$: flow of air or ethane, respectively

Materials and methods

The following reagents were used:

- $Ba(NO_3)_2$, $Sr(NO_3)_2$, $Co(NO_3)_3 \cdot 6H_2O$, $Fe(NO_3)_2 \cdot 6H_2O$, EDTA, citric acid (Sigma Aldrich®)
- $Mg(OMe)_2$ (9.34 g of Mg/L in methanol) (Epichem®)

- $\text{VO}(\text{OCH}_2\text{CH}_2\text{CH}_3)_3$ (Aldrich®).

The powders obtained were characterised by means of elemental analysis and X-ray diffraction and using the BET method.

The X-ray diffraction spectra were carried out using a system device Brücker® D5005 in the 2θ range, from 3° to 80° , a step of 0.02° , a time of 1s and a radiation of Cu $K_{\alpha 1+\alpha 2} = 1.54184 \text{ \AA}$.

The elemental composition was determined by means of ICP-OES using a spectroflame device.

Scanning electron microscopy operations were carried out using a device Hitach S-800. The transmission electron microscopy operations were carried out using a device JEOL 2010 - 200 kV.

Membrane reactor

a) Description

The oxidative dehydrogenation of ethane (DHOE) and the permeability relative to oxygen were examined using the reactor described in Figure 1.

The discs (approximately 1 mm thick) were sealed between two dense tubes of alumina (OD 12 mm, ID 8 mm) using gold as a chemically inert sealing agent.

Furthermore, the side of the wall of the disc has been extensively covered with a gold-based adhesive in order to eliminate the radial contributions of the flow of oxygen passing through the active cross-section of 0.5 cm^2 . The

welding was carried out at the beginning of the experiments by heating to a temperature of 800°C for one night. The side which is in contact with air was supplied with a constant total pressure adjusted to 120 kPa and a constant flow rate of 50 mL/min by using a mixed current of O₂ (liquid air) and N₂ (N₂ in the form of evaporated liquid air, liquid air) individually controlled by mass flow controllers (Brooks®, type 5850 TR). The side which is in contact with the fuel was supplied with a mixture of ethane (liquid air) and helium (liquid air) which is individually controlled by mass flow controllers (Brooks®, type 5850 TR) in the case of DHOE, and simply by He in the case of the permeability relative to oxygen. Two pressure sensors were installed which allow the total pressure on each surface of the reactor to be known continuously.

Figure 1 is an illustration of a high-temperature permeation membrane reactor with weld seams of gold.

b) Analysis of the gases

The reactant gases (O₂, N₂, C₂H₆) and the gases produced (H₂, CH₄, CO₂, C₂H₄, C₂H₆ and H₂O) were analysed using two chromatographs in a gaseous phase (GC), both of which are connected to a Chemstation HP in order to gather data and allow rapid analysis. The first GC (HP 5890 Series II) was equipped with a thermal conductivity detector (TCD) and a column 13X which allows the separation of O₂, N₂, CH₄, CO, and H₂O. H₂, CH₄, CO₂, C₂H₄, C₂H₆ and H₂O were eluted from a HayeSep column on the second GC (Delsi Series 200) provided with a TDC.

Furthermore, argon was introduced with the reagents in order to readily determine a possible expansion of the flow owing

to an increase in the number of moles during the process of DHOE. The balance in terms of carbon is in the order of 4%.

Gas leakages owing to a poor weld seam or incomplete densification of the membrane, should they occur, would thus be detected by following the concentration of nitrogen which allows the permselectivity in terms of O_2 to be calculated for the membrane reactor.

Example 1: Preparation of a dense $Ba_{0.5} Sr_{0.5} Co_{0.8} Fe_{0.2}$

$O_{3-\delta}$ membrane coated with particles of VMgO

*a) Preparation of the dense $Ba_{0.5} Sr_{0.5} Co_{0.8} Fe_{0.2}$
 $O_{3-\delta}$ membrane*

The perovskite powder $Ba_{0.5} Sr_{0.5} Co_{0.8} Fe_{0.2} O_{3-\delta}$ (BSCFO) was prepared using a variant of the method commonly referred to as the "citrate method". In this method, the stoichiometric quantities of $Ba(NO_3)_2$ (2.61 g; 0.5 eq), $Sr(NO_3)_2$ (2.11 g; 0.5 eq), $Co(NO_3)_2 \cdot 6H_2O$ (4.65 g; 0.8 eq) and $Fe(NO_3)_3 \cdot 6H_2O$ (4.05 mL of an aqueous solution at 0.988 mol/L; 8.76 g; 0.2 eq) (of purity > 99.5%) were dissolved in 100 mL of distilled water followed by the addition of EDTA and citric acid at a molar ratio of perovskite, EDTA and citric acid equal to 1:1.5:3.

The violet coloured solution obtained was heated to a temperature of 100°C until a gel-like material is obtained which is formed by evaporation of water after approximately three hours. The gel was then heated to 300°C for three hours. The foam obtained was then calcined at 900°C for four hours in air in order to lead to the production of a perovskite powder. This powder was then moulded in a mortar. Bare membrane discs were compressed at a pressure of 140 MPa for one minute. The densification of the discs was then carried

out by means of sintering at 1150°C for eight hours. The thickness of the membrane is from 1mm +/- 0.1 and the surface-area is 0.5 cm².

b) Preparation and deposit of the catalyst particles VMgO

The particles based on mixed magnesium oxide doped using 10% by weight of vanadium were prepared from molecular precursors using the sol-gel method. This allows thin layers to be deposited which have high levels of purity and specific surface-area.

Figure 2 illustrates the specific surface-area of VMgO in accordance with temperature.

The magnesium alkoxide used ($\text{Mg}(\text{OMe})_2$) (9.34g of Mg/L of methanol) was prepared by Epichem® and vanadium alkoxide ($\text{VO}(\text{OCH}_2\text{CH}_3)_3$) (208.8g of V/L of methanol) is a solution marketed by Aldrich®. The two solutions were mixed so as to obtain a solution in which vanadium represents 10% by weight.

A few drops of the solution obtained, corresponding to a quantity of approximately 1 mg of VMgO were then deposited on the surface of the membrane using the technique of "spin-coating". The membrane was then placed directly in the reactor at 850°C. Particles of VMgO are thus obtained of 2µm in size, which are distributed according to a density of $[5 \cdot 10^6 \text{ part/cm}^2 \text{ (to within 10\%)}]$ and which cover approximately 40% of the surface-area of the membrane (to within 10%).

Figure 12 illustrates the surface of the membrane using electron microscopy (Scale: 10µm/cm).

c) Features of the powders

The formation of a pure perovskite structure was verified by means of X-ray diffraction.

The elemental composition was determined by ICP-OES (spectroflame) by analysing a sample dissolved by heating from 250° to 300°C in a mixture of H₂SO₄ and HNO₃ after calcination.

Sample	Ba/at.-%	Sr/at.-%	Co/at.-%	Fe/at.-%
Batch 1	24.9	25.1	39.9	10.1
Batch 2	25.0	25.0	40.0	10.0

Table 1: Cationic Composition of the BSCFO powder.

Results of the ICP-OES analysis.

Table 1 illustrates that this method ensures a very good reproducibility of the composition of the powder.

Example 2: Preparation of a dense BSCFO membrane coated with particles of catalyst MgO

The dense Ba_{0.5} Sr_{0.5} Co_{0.8} Fe_{0.2} O_{3-δ} membrane is prepared in accordance with that of example 1.

The particles based on magnesium oxide were prepared using the sol-gel technique from a methanol-containing solution of Mg(OMe)₂ (Epichem®) at 9.34 g/L.

1 mg of the colloidal suspension obtained was deposited on the surface of the membrane using the technique of spin-coating. The membrane was then placed directly in the reactor at 850°C.

Particles of MgO are thus obtained of 2 μ m in size, which are distributed according to a density of $[5 \cdot 10^6 \text{ part/cm}^2 \text{ (to within 10\%)}]$ and which cover approximately 40% of the surface-area of the membrane.

Example 3: Dense BSCFO membrane coated with particles of Pd catalyst

The dense membrane used is in accordance with that described in example 1.

The Pd particles were deposited at the surface of the dense membrane in accordance with the technique of laser vaporisation. The laser ablation of a Pd bar of 99.99% purity (Goodfellow) was carried out using an Nd:YAG(BMI®) laser which is pulsed at 30 Hz and doubled in terms of frequency (532 nm).

The quantity of material deposited at the surface of the membrane was determined using a quartz microbalance (INFICON®).

Particles of Pd are thus obtained which have a mean size of 2-3 nm, which are distributed according to a density of $4 \cdot 10^{12} \text{ part/cm}^2$, and which cover approximately 8% of the membrane (to within 2%).

Figure 13 is an illustration of the surface of the membrane coated with dispersed particles of Pd using electron microscopy.

Example 4: Determination of the speed of oxygen permeation

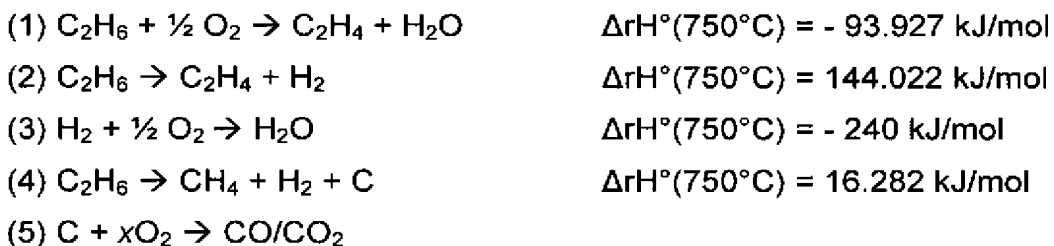
Application of the membrane according to example 2 to the separation of oxygen.

The stability of the membrane of example 2 with regard to permeation of oxygen was examined by continuously recording the permeation flow at a constant partial pressure gradient of oxygen for 30 hours

As described in Figure 3, a stable permeation flow of oxygen of 2.4 mL.min.cm² is obtained at 725°C. These results represent a flow which is 1.5 times greater than that corresponding to the bare membrane.

Example 5: Application of the membrane to the oxidative dehydrogenation of ethane

a) Main possible reactions



b) Calculation

Experiments for oxidation of ethane and propane in membrane reactors were carried out at different temperatures 700-825°C, different flow rates C₂H₆/He (35-65-100 mL/min) and at partial pressures of C₂H₆ of 0.25-0.44-0.55 atm at the side of the membrane which is in contact with ethane (oxidation zone).

All the calculations were carried out based on the partial pressure P_x , the total flow $F^\circ_{\text{ethane side}}$, the expansion coefficient of the flow C_{exp} and the surface-area of the

active membrane S (based on the assumption of behaviour in accordance with the law for an ideal gas).

c) Effect of temperature

Figure 4 illustrates the yield of ethylene in accordance with temperature. This increases up to 775°C in order to reach 75% (conversion=87%, selectivity=87%). At 700°C, the low yield (35%) is a result of a poor conversion (<40%) since the selectivity is greater than 90% as indicated in Figure 5. However, at 825°C, the low yield (50%) is a result of a reduction of the selectivity of ethylene (<60%) in favour of carbon monoxide.

d) Effect of the partial pressure of ethane

Figure 6 illustrates the yield of ethylene in accordance with the partial pressure of ethane at the "optimum" temperature of 775°C. No variation of the yield in accordance with the partial pressure of ethane is seen. The selectivity and the conversion are also relatively constant.

Example 6: Application of membranes according to examples 1 to 3 to the oxidative dehydrogenation of ethane

Figure 7 illustrates the ethylene yield of the membrane according to example 3 in accordance with temperature. This increases up to 775°C, in order to reach a maximum of 75%, which corresponds to a conversion rate of 87% and a selectivity rate of 87%.

Figure 8 compares the ethane yields at different temperatures for dense membranes which are simple, that is to say, bare, and which are partially coated with dispersed particles based on magnesium oxide (examples 1 and 2) or Pd (example 3).

The presence of dispersed particles of Pd based on magnesium oxide allows a specific yield to be achieved at a temperature which is on average 75°C lower than that of dense membranes.

The presence of dispersed particles of Pd or based on magnesium oxide in particular allows yields to be obtained which are greater than those of dense membranes (Y=75%, C=87%, S=87%) at a specific temperature.

The yields obtained with particles of catalyst based on magnesium oxide and Pd are comparable.

Example 7: Application of the membrane of example 2 to the oxidative dehydrogenation of propane

a) Effect of temperature

Figure 9 indicates the yields of propylene, ethylene and total alkene (i.e. propylene and ethylene) in accordance with temperature. For propylene, a maximum yield of 19% (S=27.4%, C=69.7%) is obtained at 725°C. This low yield is principally the result of a secondary cracking reaction of the propane under the operating temperature conditions of the membrane. The main product obtained is ethylene (S=47.2% at 725°C), which is a very valuable product.

b) Effect of the partial pressure of propane

Figure 10 illustrates the yield of propylene and ethylene at 700°C in accordance with the partial pressure of propane.

The yield in terms of ethylene and propylene remains constant for a partial pressure in terms of propane of between 0.15 and 0.50 bar.

Example 8: Application of the membrane of example 3 to the oxidative dehydrogenation of propane

Figure 11 indicates the yields of propylene, ethylene and alkene (propylene and ethylene) in accordance with the temperature. For propylene, a maximum yield of 18.2% (S=25.9%, C=70.1%) is obtained at 725°C.

Example 9: Comparative study of the membranes of examples 1, 2 and 3 and the membranes of the prior art

The permeation flows of oxygen of various membranes were determined using the reactor described in Figure 1 under the conditions indicated in the above section "Material and methods".

These results indicate that the modification of one of the surfaces of the dense membrane using particles of noble metals or based on magnesium oxide allows the permeation flow of oxygen to be significantly increased.

Membrane	T (°C)	Flow of oxygen	Reference
Without surface-area modification	800	Mol cm ⁻² s ⁻¹ 6.69 E-07	[1]
Ba _{0.5} Sr _{0.5} Co _{0.8} Fe _{0.2} O _x	800	3.12 E-07	[2]
BaBi _{0.4} Co _{0.2} Fe _{0.4} O _{3-δ}	800	1.12 E-07	[3]
La _{0.2} Sr _{0.8} Co _{0.2} Fe _{0.8} O _{3-δ}	800	1.86 E-08	[4]
La _{0.6} Sr _{0.4} Co _{0.2} Fe _{0.8} O _{3-δ}	800	5.20 E-08	[5]
La _{0.5} Sr _{0.4} Co _{0.8} Fe _{0.2} O _{3-δ}	800	4.80 E-07	[1]
SrCo _{0.8} Fe _{0.2} O _{3-δ}	800	3.35 E-07	[7]
BaCo _{0.4} Fe _{0.4} Zr _{0.2} O ₃₋₁	800	1.58 E-08	[8]
Bi ₂ V _{0.8} Mn _{0.2} O _{5.3}	800		

With surface-area modification $Ba_{0.5}Sr_{0.5}Co_{0.8}Fe_{0.2}O_{x+MgO}$	800	1.00 E-06	Example 2
Ibid + Pd	800	1.12 E-06	Example 3
Ibid + VMgO	800	1.27 E-06	Example 1

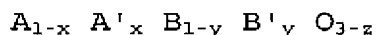
Table 2

1. Shao, Z., et al., Investigation of the permeation behavior and stability of a $Ba_{0.5}Sr_{0.5}Co_{0.8}Fe_{0.2}O_{3-\delta}$ oxygen membrane. *Journal of Membrane Science*, 2000. 172(1-2): p. 177-188.
2. Shao, Z., et al., Synthesis and oxygen permeation study of novel perovskite-type $Ba_{1-x}Bi_xCo_{0.2}Fe_{0.8-x}O_{3-\delta}$ ceramic membranes. *J. Membr. Sci.*, 2000. 164(1-2): p. 167-176.
3. Li, S.G., et al., Comparison of oxygen permeability and stability of perovskite type $La(0.2)A(0.8)Co(0.2)Fe(0.8)O(3-\delta)$ ($A = Sr, Ba, Ca$) membranes. *Industrial and Engineering Chemistry Research*, 1999. 38(8): p. 2963-2972.
4. Lee, S., et al., Oxygen-permeating property of $LaSrBFeO_3$ ($B=Co, Ga$) perovskite membrane surface-modified by $LaSrCoO_3$. *Solid State Ionics*, 2003. 158(3-4): p. 287-296.
5. Teraoka, Y., et al., Catalytic effects in oxygen permeation through mixed-conductive LSCF perovskite membranes. *Solid State Ionics*, 2002. 152-153: p. 681-687.
6. Qiu, L., Lee T. H., Oxygen permeation studies of $SrCo_{0.8}Fe_{0.2}O_{3-\delta}$. *Solid State Ionics*, 1995. 76: p. 321-329.
7. Tong, J., et al., Investigation of ideal zirconium-doped perovskite-type ceramic membrane materials for oxygen separation. *Journal of Membrane Science*, 2002. 203(1-2): p. 175-189.

8. Yang, Y.L., Qiu L. Jacobson, *Manganese doped bismuth vanadate solid electrolytes: oxygen permeation in $\text{Bi}_2\text{V}_{0.8}\text{Mn}_{0.2}\text{O}_{5.3}$* . *Journal of Materials Chemistry*, 1997. 7: p. 937-941.

CLAIMS

1. Oxygen conducting membrane comprising a mixed conducting dense membrane of multimetal oxide, one of the surfaces of which is covered with dispersed particles based on magnesium oxide or noble metals.
2. Membrane according to claim 1, in which the particles based on magnesium oxide and/or noble metals have a diameter of between 5 and 3000 nm.
3. Membrane according to either claim 1 or claim 2, in which the dense mixed conducting membrane of multimetal oxide has a perovskite structure.
4. Membrane according to any one of claims 1 to 3, in which the dense mixed conducting layer comprises one or more multimetal oxides which comply with the general formula :



where

A and A', which may be the same or different, each represent a metal ion or an alkaline-earth metal or a metal which is selected from the lanthanide series;

B and B', which may be the same or different, each represent a metal ion and/or a mixture of metal ions in which the metal is selected from the transition metals;

$$0 \leq x \leq 1;$$

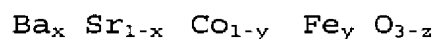
$$0 \leq y \leq 1;$$

z is a number which renders the charge of the compound neutral and which defines the oxygen deficiency.

5. Membrane according to claim 4, in which A and A', which may be the same or different, represent La, Ca, Sr, and/or Ba.

6. Membrane according to either claim 4 or claim 5, in which B and B', which may be the same or different, are selected from Cr, Mn, Fe, Co, Ni and/or Cu.

7. Membrane according to any one of claims 4 to 6, in which the dense mixed conducting layer comprises multimetal oxide having the formula:



where x, y and z are as defined in claim 6.

8. Membrane according to claim 7, in which the multimetal oxide comprises $\text{Ba}_{0.5} \text{Sr}_{0.5} \text{Co}_{0.8} \text{Fe}_{0.2} \text{O}_{3-z}$.

9. Membrane according to any one of the preceding claims, in which the dense mixed conducting membrane of multimetal oxide has a thickness of between 0.5 and 10 mm.

10. Membrane according to any one of the preceding claims, in which the particles based on magnesium oxide or noble metals represent from 0.01 to 0.1% by weight of the dense membrane.

11. Membrane according to any one of the preceding claims, in which the particles are based on magnesium oxide.

12. Membrane according to claim 11, in which the particles based on magnesium oxide are doped using vanadium.

13. Membrane according to any one of claims 1 to 10, in which the particles are particles of noble metals or alloys thereof.

14. Membrane according to claim 13, in which the noble metals are selected from Pd, Pt, Rh, Ag, Au, Ru and Ir.

15. Method for preparing oxygen conducting membranes as defined according to claims 1 to 12, comprising the steps consisting in:

- a. providing a dense mixed conduction membrane as defined in claims 1 and 3 to 9;
- b. preparing a colloidal suspension based on magnesium oxide in an organic solvent;
- c. placing the suspension obtained in contact with the dense mixed conducting membrane; and
- d. calcining the membrane obtained.

16. Method for preparing oxygen conducting membranes as defined according to claims 1 to 10 and 13 to 14, comprising the steps consisting in:

- a. providing a dense membrane of multimetal oxide as defined in claims 1 and 3 to 9;
- b. depositing the particles of noble metals or alloys thereof by means of laser vaporisation.

17. Oxygen conducting membrane which can be obtained using the method according to either claim 15 or claim 16.

18. Membrane reactor comprising an oxidation zone and a reduction zone which are separated by means of an oxygen conducting membrane as defined in any one of claims 1 to 14 or 17.

19. Membrane reactor according to claim 18, in which the oxidation zone is in contact with the surface of the membrane

coated with dispersed particles based on magnesium oxide or noble metals.

20. Method for oxidising a reactant gas comprising:

- i) using a membrane reactor according to claims 18 or 19;
- ii) introducing the reactant gas into the oxidation zone;
- iii) introducing the gas containing oxygen into the reduction zone;
- iv) heating the membrane which separates the oxidation and reduction zones to an operating temperature.

21. Method according to claim 20, in which the reactant gas is a light hydrocarbon which is oxidised into alkene.

22. Method according to claim 21, in which the light hydrocarbon is ethane which is oxidised into ethylene.

23. Use of a membrane reactor according to claims 18 or 19 to recover oxygen from a gaseous mixture containing oxygen.

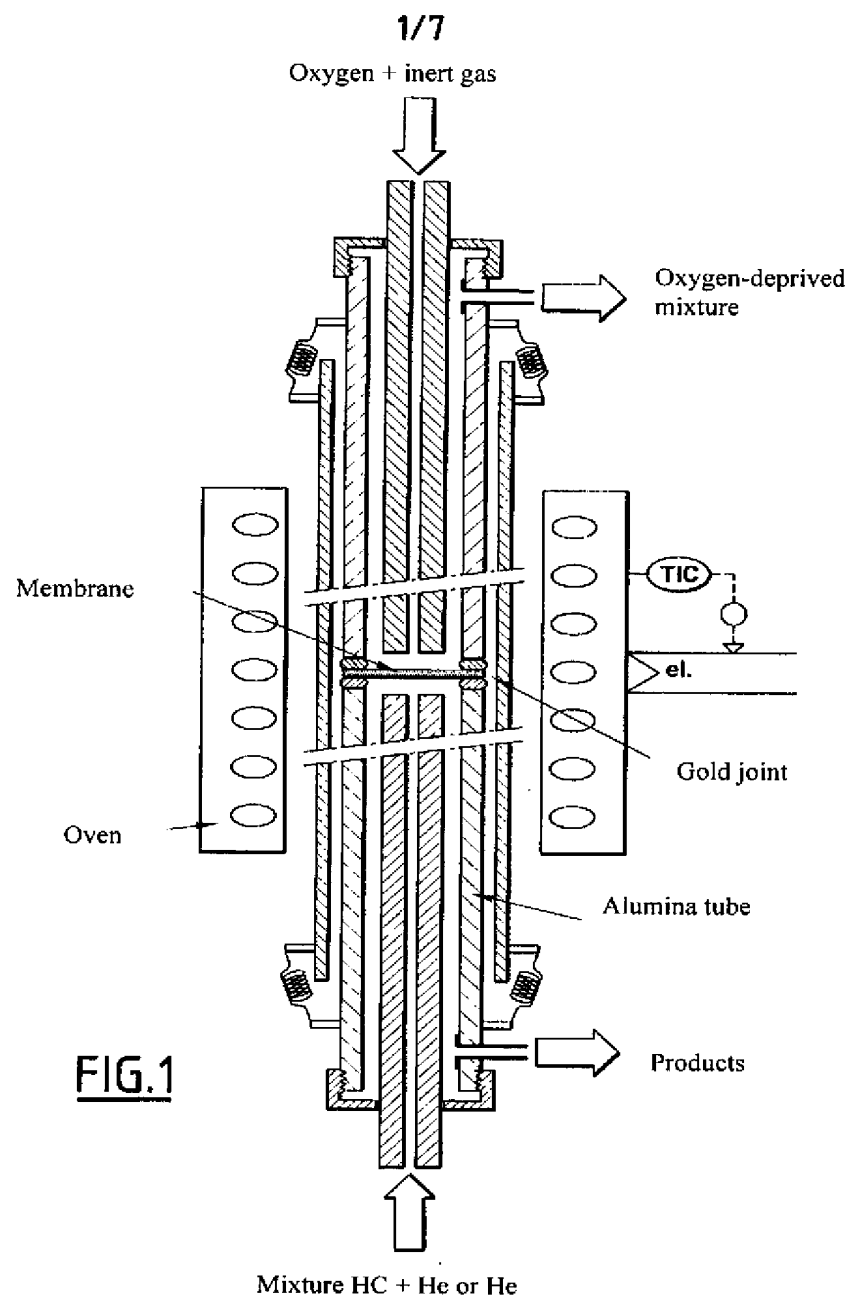


FIG.1

2/7

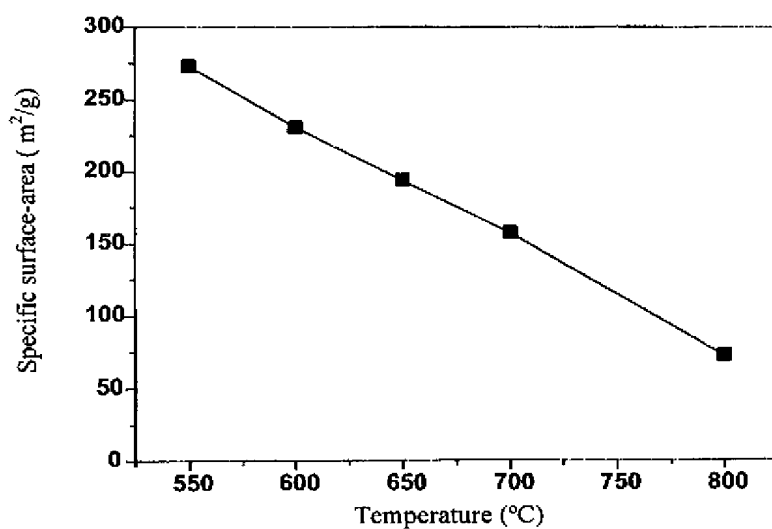


FIG.2

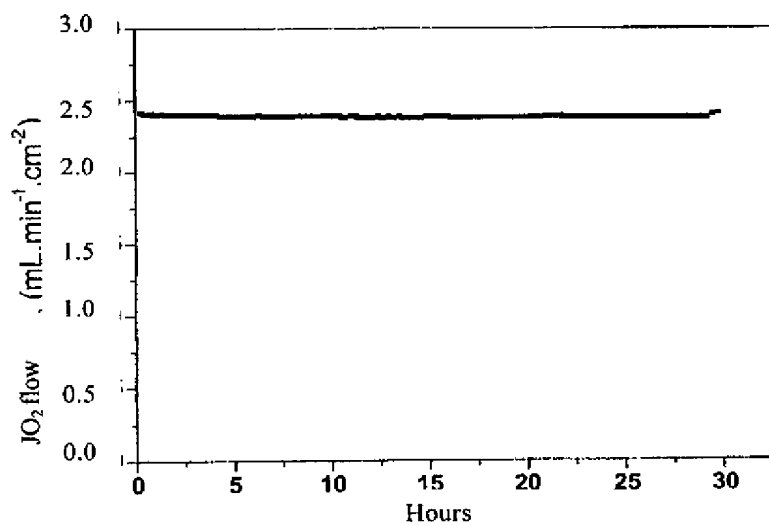


FIG.3

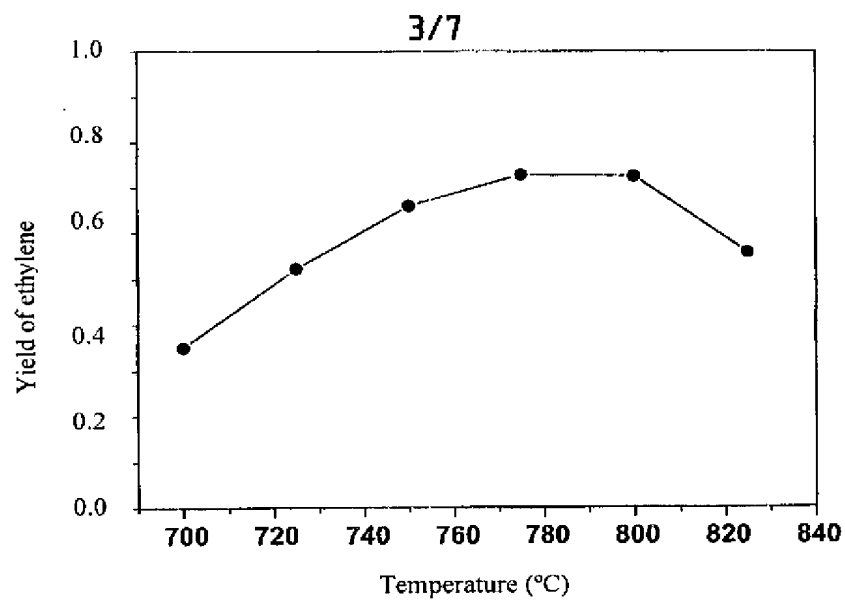


FIG.4

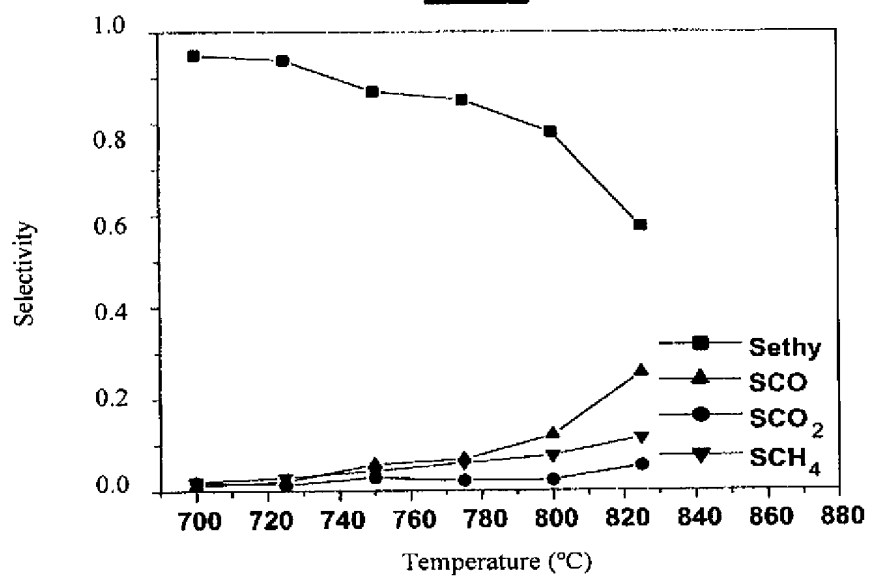


FIG.5

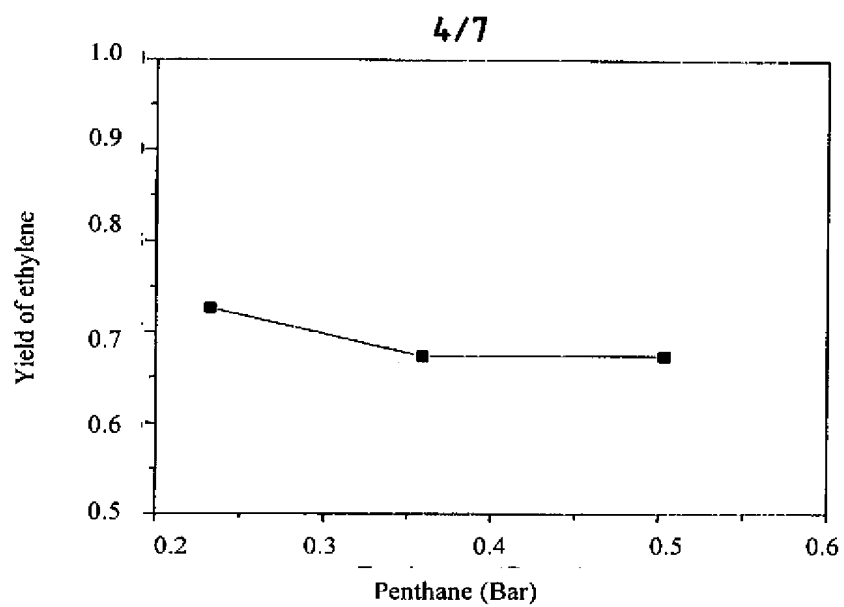


FIG.6

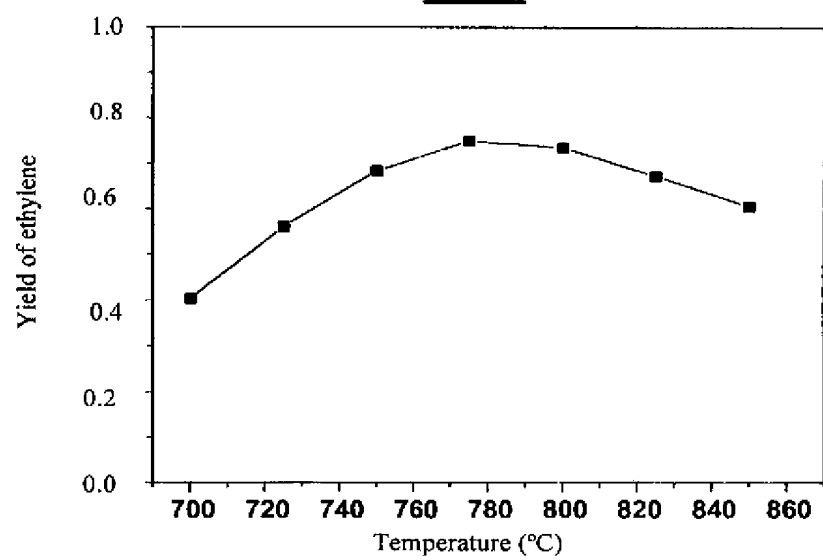


FIG.7

5/7

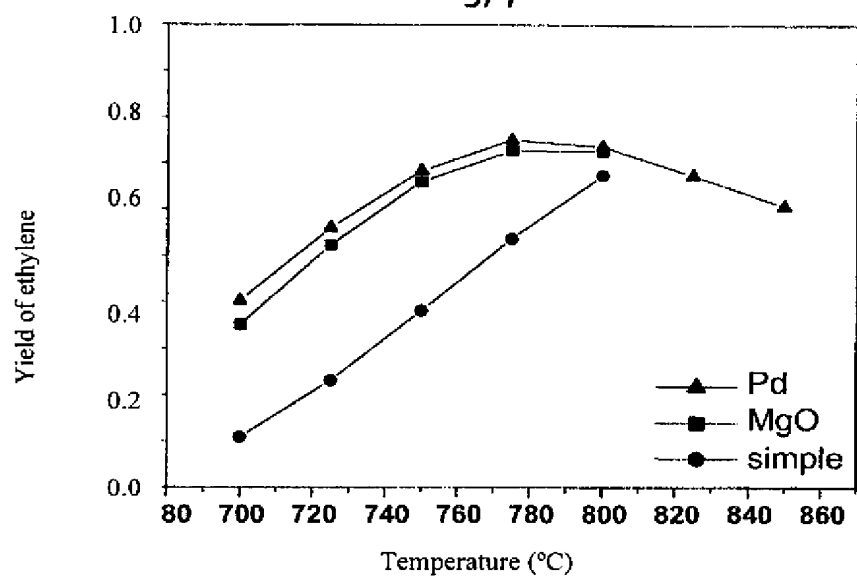


FIG.8

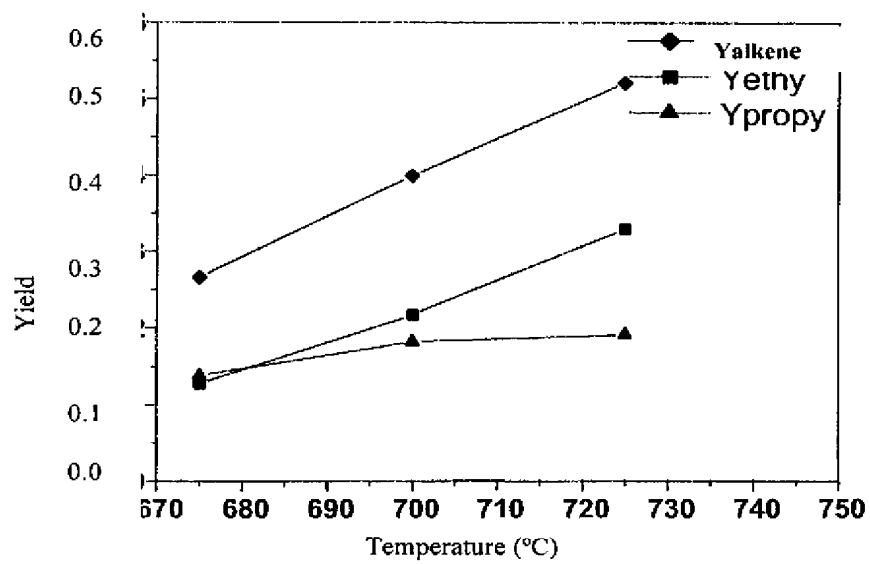


FIG.9

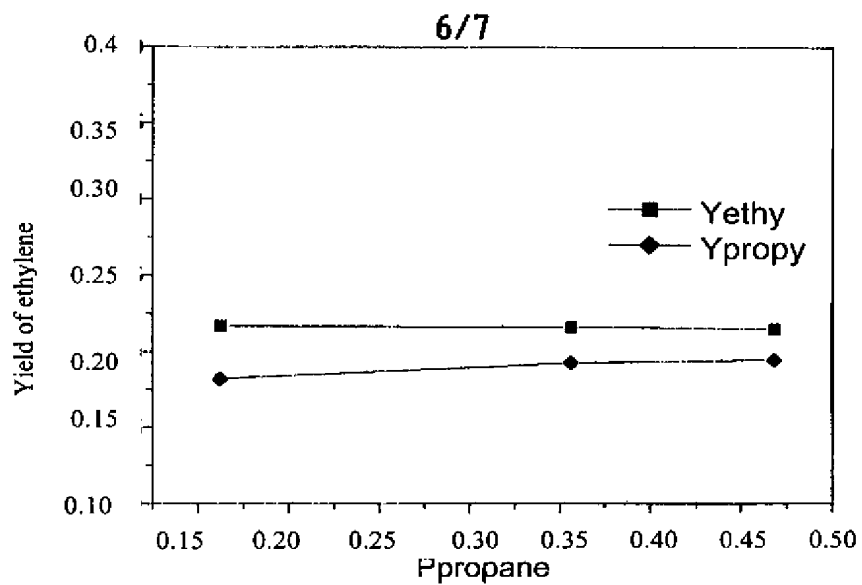


FIG.10

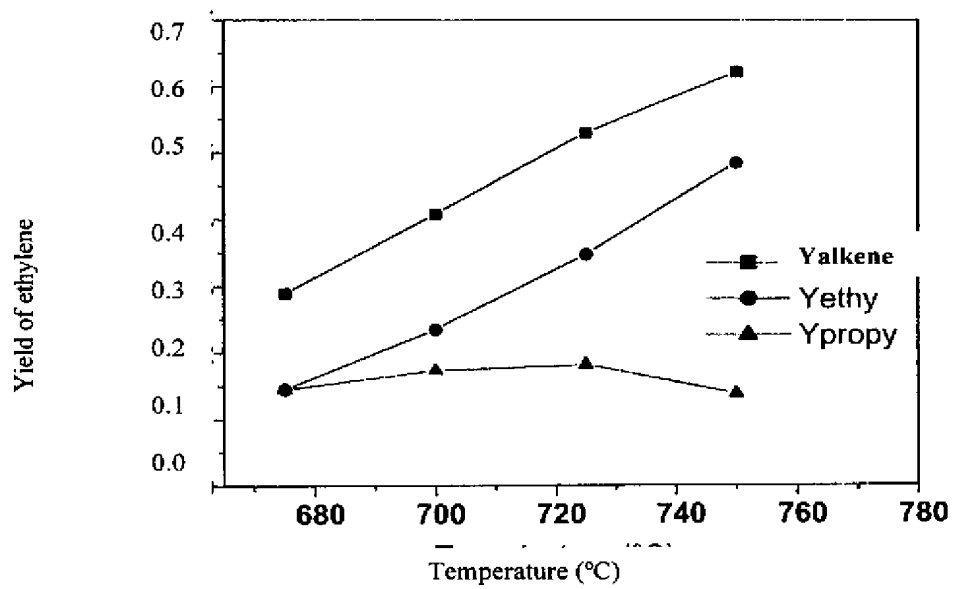


FIG.11

7/7



FIG.12

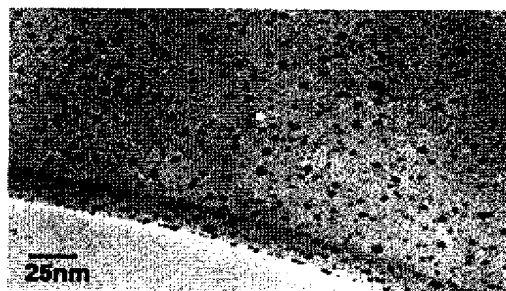


FIG.13